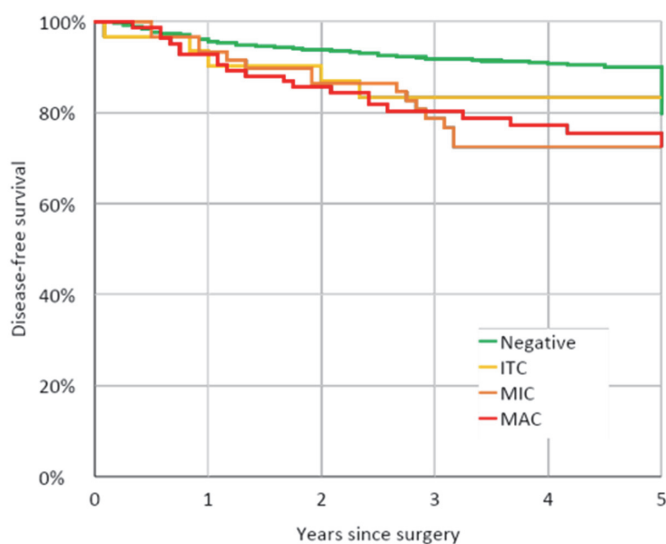


Disease-free survival: All patients by largest type of metastasis in LN (N = 969)



Number at risk (cumulative number of events)

	0	1	2	3	4	5
Negative	795 (0)	763 (31)	721 (49)	501 (62)	364 (66)	223 (69)
ITC	31 (0)	29 (2)	27 (3)	19 (5)	15 (5)	12 (5)
MIC	59 (0)	55 (4)	51 (8)	38 (12)	28 (15)	23 (15)
MAC	84 (0)	78 (6)	71 (12)	54 (16)	45 (18)	29 (19)

Abstract 898 Figure 1

Abstract 898 Table 2 Univariate analysis of factors associated with disease-free survival (N = 969)

Predictor	Category	n	HR (95% CI)	p-value
Surgical approach	Open	575	Ref.	
	Robotic	195	1.21 (0.74; 1.97)	0.439
	Laparoscopic	141	1.51 (0.93; 2.45)	0.097
	Combined	58	1.06 (0.48; 2.31)	0.888
Tumour diameter	< 0.5 cm	73	Ref.	
	0.5–1.99 cm	424	1.67 (0.51; 5.47)	0.399
	2–3.99 cm	376	3.98 (1.25; 12.69)	0.019
	≥ 4 cm	96	6.35 (1.91; 21.13)	0.003
LVSI	No	316	Ref.	
	Yes	351	2.31 (1.47; 3.63)	< 0.001
Tumour histotype	Squamous	605	Ref.	
	Adenocarc.	287	1.13 (0.75; 1.71)	0.554
	Adenosquamous	50	1.38 (0.66; 2.89)	0.385
	Other	27	3.03 (1.45; 6.31)	0.003
Grade	1	149	Ref.	
	2	406	2.08 (1.02; 4.22)	0.044
	3	246	3.35 (1.64; 6.85)	< 0.001
Largest type of metastasis in LN	Negative	795	Ref.	
	ITC	31	1.67 (0.68; 4.14)	0.264
	MIC	59	2.55 (1.47; 4.43)	< 0.001
	MAC	84	2.36 (1.44; 3.87)	< 0.001
Largest type of metastasis in LN	Negative	795	Ref.	
	ITC	31	1.67 (0.68; 4.14)	0.264
	MIC+MAC	143	2.44 (1.63; 3.64)	< 0.001

Result(s)* Out of 969 included patients with at least 1 SLN detected, 174 (18%) had positive LN (table 1). Maximal tumour diameter >2cm, positive LVSI, grade ≥ 2, uncommon histological type (neuroendocrine, sarcoma, etc.) and macro-metastasis (MAC) or MIC in LN were factors associated with significantly decreased five-years disease free survival (DFS) (table 2). MAC, MIC or ITC was the largest LN metastasis in 84 (9%), 59 (6%) and 31 (3%) cases respectively. Adjuvant (chemo)radiation was administered in 89%, 85% and 58% of patients with MAC, MIC and ITC. DFS reached 75%, 73% and 83% in patients with MAC, MIC and ITC compared with 90% in the N0 patients. Patients with MAC and MIC had significantly decreased DFS than those with N0 disease (HR=2.36 and 2.55).

Conclusion* Early-stage cervical cancer patients with MIC in pelvic LN have significantly decreased DFS. Their management should follow the same principles as in patients with MAC.

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PHASE 1B TRIAL OF FIRST-LINE BINTRAFUSP ALFA, A BIFUNCTIONAL FUSION PROTEIN TARGETING TGF-β AND PD-L1, PLUS CHEMOTHERAPY WITH OR WITHOUT BEVACIZUMAB IN CERVICAL CANCER

¹A Oaknin*, ²M Gil-Martin, ³E Diver, ⁴G Jehl, ⁴SA Gleicher, ⁵S Chaudhary, ⁵L Ojalvo, ⁶K Hasegawa. ¹Vall d'Hebron University Hospital, Vall d'Hebron Institute of Oncology (VHIO), Barcelona, Spain; ²ICO Hospital Duran i Reynals, Barcelona, Spain; ³Stanford Cancer Institute, Stanford, CA, USA; ⁴Merck KGaA, Darmstadt, Germany; ⁵EMD Serono Research and Development Institute, Inc., Billerica, MA, USA; ⁶Saitama Medical University International Medical Center, Hidaki-shi, Saitama-ken, Japan

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Introduction/Background* Platinum-containing chemotherapy ± bevacizumab is standard-of-care for recurrent/metastatic/persistent (R/M/P) cervical cancer (CC). Anti-PD-(L)1 therapy has benefit in some patients who progress after first-line (1L) therapy; 1L efficacy is unknown. HPV infection, implicated in >95% of CCs, is linked to TGF-β upregulation. Bintrafusp alfa is a first-in-class bifunctional fusion protein composed of the extracellular domain of the TGF-βRII receptor (a TGF-β ‘trap’) fused to a human IgG1 mAb blocking PD-L1. Promising activity was observed in patients with recurrent, platinum-experienced CC (response rate 28.2%). We report data from a phase 1b trial evaluating safety of 1L bintrafusp alfa + chemotherapy ± bevacizumab (INTR@PID 046; NCT04551950).

Methodology Patients with R/M/P CC who had not received prior systemic therapy were eligible for cohort 1. They received bintrafusp alfa 2400mg q3w plus cisplatin 50mg/m² or carboplatin AUC5, paclitaxel 175mg/m² with (cohort 1A)/without (cohort 1B) bevacizumab 15mg/kg until disease progression, death, unacceptable toxicity, or withdrawal. Primary endpoints: occurrence of predefined dose-limiting toxicities (DLT) ≤4 weeks from treatment start; adverse event occurrence. Target recruitment was 8 patients/cohort, with safety assessments when 3 and 8 patients had completed the DLT period.

Result(s)* As of May 4, 2021, 8 and 9 patients in cohorts 1A and 1B had received therapy for a median of 10.6 and 9.0 weeks. All patients had completed the DLT period and remained on therapy. Two non-bintrafusp alfa-related DLTs were observed in cohort 1B (grade 4 amylase elevation, grade 3 menorrhagia); neither led to treatment discontinuation. Any-grade treatment-related adverse events (TRAEs) occurred in 62.5% and 100% of patients in cohorts 1A and 1B. Grade 3 TRAEs occurred in 3 and 2 patients (cohort 1A: anemia [n=2], lipase increase, decreased neutrophil count, maculopapular rash [n=1 each]; cohort 1B: anemia, rectal hemorrhage, vaginal bleeding [n=1 each]); 1 patient in cohort 1B had grade 4 anemia. No treatment-related deaths occurred. Preliminary efficacy based on short follow-up showed 3 and 2 tumor responses (2 and 1 pending confirmation) in cohorts 1A and 1B.

Conclusion* No new safety signals were observed with 1L bintrafusp alfa + chemotherapy ± bevacizumab in patients with R/M/P CC. Further studies are warranted.

919 ULTRASOUND ROLE IN STAGING OF CANCER CERVIX

A Elagwany*. Alexandria university , Obg , Alexandria, Egypt

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Introduction/Background* Cancer cervix is common in developing countries due to limited adoption of screening programs . It is far less common in Developed countries due to availability of national screening program and governmental health insurance system . Cancer cervix in developing countries usually present in stage 1b and beyond . MRI and EUA is usually used for staging before surgery .

Methodology Recently, due to advanced technologies in ultrasound , we can now stage cancer cervix accurately and replace MRI and EUA . We are trying here to spot the lights over this with a pictorial illustration of different stages. The accuracy of vaginal sonography for the evaluating cancer cervix is

comparable to that of MRI and even better for local staging in identifying tissue planes

Result(s)* Ultrasound can be used in cancer cervix to assess the topography regarding exophytic versus endophytic tumor, The tumor size measured in three diameters and the distance between the tumor and the internal cervical os .the pericervical fascia which is the paracervix at he level of the cervix and the paracolpos at the level of the vagina is assessed . Thence, the extent of the radical procedure (parametrectomy) can be planned.

The tumor is usually hypoechoic in cases of squamous cell carcinoma and hyperechoic in adenocarcinoma . His is important in differentiating large bulky endocervical tumors (with regular outline) (stage 1) from those with parametrial invasion with irregular outline (stage 2 b) . The vaginal extension is evaluated by the thickening or masses of the vaginal walls (stage 2a) along with assessing the paracolpos in the same manner as before (stage 2b). Ureteral dilatation is common in parametrial infiltration and is seen .

The spread into the urinary bladder and/or rectum (stage 4) can be determined and the ultrasound for the bladder involvement is better than does cystoscopy, as this can only show bullous mucosal edema or mucosal lesion but not the wall affection that can be seen by ultrasound . The assessment of both is based on assessing her muscle layer and the related fascia.

Conclusion* Ultrasound is comparable to mri in local staging of cancer cervix especially for minor changes .

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A LARGE, MULTICENTER, RETROSPECTIVE STUDY ON EFFICACY AND SAFETY OF STEREOTACTIC BODY RADIOTHERAPY IN OLIGOMETASTATIC CERVICAL CANCER (MITO-RT/RAD)

¹G Macchia, ²M Campitelli, ¹P Bonome, ³C Lalischia, ⁴A Fodor, ⁵L Draghini, ⁶P Gentile, ⁷GR D'agostino, ⁸V Balcet, ⁹A Raguso, ¹⁰E Ippolito, ^{11,12}M Ferioli, ¹³L Vicenzi, ¹⁴S Borghesi, ²P Mitidieri, ¹⁵VDI Cataldo, ¹⁶E Perrucci, ¹⁷S Pignata, ¹⁸G Scambia, ¹⁹G Ferrandina*. ¹Gemelli Molise Hospital – Università Cattolica del Sacro Cuore, Radiation Oncology Unit, Campobasso, Italy; ²Fondazione Policlinico Universitario A. Gemelli, IRCCS, UOC di Radioterapia, Dipartimento di Scienze Radiologiche, Radioterapiche ed Ematologiche,, Rome, Italy; ³University of Pisa, Department of Translational Medicine, Division of Radiation Oncology, Pisa, Italy; ⁴IRCCS San Raffaele Scientific Institute, Department of Radiation Oncology, Milan, Italy; ⁵S. Maria Hospital, Radiation Oncology Centre, Terni, Italy; ⁶UPMC Hillman Cancer Center San Pietro FBF, Radiation Oncology Unit, Rome, Italy; ⁷Humanitas Clinical and Research Hospital, IRCCS, Department of Radiotherapy and Radiosurgery, Rozzano, Italy; ⁸Nuovo Ospedale degli Infermi, UOC Radioterapia, Biella, Italy; ⁹Fondazione ‘Casa Sollievo della Sofferenza’, IRCCS, UOC Radioterapia, S. Giovanni Rotondo (FG); ¹⁰Campus Bio-Medico University, Department of Radiation Oncology, Rome, Italy; ¹¹Radiation Oncology, IRCCS Azienda Ospedaliero-Universitaria di Bologna, Bologna; ¹²Department of Experimental, Diagnostic and Specialty Medicine – DIMES, Alma Mater Studiorum University of Bologna; ¹³Azienda Ospedaliera Universitaria Ospedali Riuniti, Radiation Oncology Unit, , Ancona, Italy; ¹⁴Azienda USL Toscana Sud Est, Radiation Oncology Unit of Arezzo-Valdarno, Arezzo, Italy; ¹⁵University of Florence, Radiation Oncology Unit, Oncology Department, Firenze, Italy; ¹⁶Perugia General Hospital, Italy, Radiation Oncology Section, Perugia, Italy; ¹⁷Istituto Nazionale Tumori di Napoli, Fondazione Pascale IRCCS, Naples, Italy; ¹⁸Fondazione Policlinico Universitario A. Gemelli, IRCCS, UOC Ginecologia Oncologica, Dipartimento per la salute della Donna e del Bambino e della Salute Pubblica, Roma, Italy; ¹⁹Fondazione Policlinico Universitario A. Gemelli, IRCCS, UOC Ginecologia Oncologica, Dipartimento per la salute della Donna e del Bambino e della Salute Pubblica, Roma, Italy

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Introduction/Background* Data supporting stereotactic body radiotherapy (SBRT) for oligometastatic gynecological cancer patients are increasing, but stereotactic treatments have not